

WHAT IS CLAIMED IS:

1. A method method for treating a thromboembolic disorder, comprising: administering, to a host in need of such treatment, a therapeutically effective amount of:

5 (a) a first therapeutic agent which is 1-(3'-aminobenzisoxazol-5'-yl)-3-trifluoromethyl-5-[[4-[(2'-dimethylaminomethyl)imidazol-1'-yl]-2-fluorophenyl]aminocarbonyl]pyrazole-hydrochloric acid salt (Compound A) or a pharmaceutically acceptable salt
10 form thereof; and,

(b) a second therapeutic agent which is Clopidogrel or a pharmaceutically acceptable salt form thereof.

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2. A method according to Claim 1, wherein at least one of the first and second therapeutic agents is administered in a sub-therapeutic dosage.

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3. A method according to Claim 2, wherein both the first and second therapeutic agents are administered in sub-therapeutic dosages.

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4. A method according to Claim 1, wherein the first and second therapeutic agents are administered simultaneously.

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5. A method according to Claim 1, wherein the first and second therapeutic agents are administered sequentially.

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6. A method according to Claim 1, wherein the method further comprises: administering, to a host in need of such treatment, a therapeutically effective amount of:

5 (c) a third therapeutic agent selected from other anti-coagulant or coagulation inhibitory agents, anti-platelet or platelet inhibitory agents, thrombin inhibitors, thrombolytic agents, fibrinolytic agents, anti-arrythmic agents, and cholesterol/lipid lowering
10 agents.

7. A method according to Claim 6, wherein the third therapeutic agent is selected from aspirin and
15 pravastatin.

8. A method according to Claim 7, wherein the third therapeutic agent is aspirin.
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9. A method according to Claim 6, wherein the third therapeutic agent is administered simultaneously with the first and second therapeutic agents.
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10. A method according to Claim 6, wherein the third therapeutic agent is administered in a sub-therapeutic dosage.
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11. A method according to Claim 1, wherein the thromboembolic disorder is selected from an arterial cardiovascular thromboembolic disorder, a venous
35 cardiovascular thromboembolic disorder, an arterial

cerebrovascular thromboembolic disorder, and a venous cerebrovascular thromboembolic disorder.

5 12. A method according to Claim 11, wherein the thromboembolic disorder is selected from unstable angina, first myocardial infarction, recurrent myocardial infarction, ischemic sudden death, transient ischemic attack, stroke, atherosclerosis, venous thrombosis, deep
10 vein thrombosis, thrombophlebitis, arterial embolism, coronary arterial thrombosis, cerebral arterial thrombosis, cerebral embolism, kidney embolism, pulmonary embolism, and thrombosis resulting from (a) prosthetic valves or other implants, (b) indwelling catheters, (c)
15 stents, (d) cardiopulmonary bypass, (e) hemodialysis, or (f) other procedures in which blood is exposed to an artificial surface that promotes thrombosis.

20 13. A method according to Claim 12, wherein the thromboembolic disorder is selected from unstable angina, first myocardial infarction, recurrent myocardial infarction, ischemic sudden death, transient ischemic attack, and stroke.

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 14. A pharmaceutical composition, comprising:
 (a) a first therapeutic agent which is 1-(3'-aminobenzisoxazol-5'-yl)-3-trifluoromethyl-5-[[4-[(2'-
30 dimethylaminomethyl)imidazol-1'-yl]-2-fluorophenyl]aminocarbonyl]pyrazole-hydrochloric acid salt (Compound A) or a pharmaceutically acceptable salt form thereof;

(b) a second therapeutic agent which is Clopidogrel or a pharmaceutically acceptable salt form thereof; and,

(c) a pharmaceutically acceptable carrier.

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15. A pharmaceutical composition according to Claim 14, wherein at least one of the first and second therapeutic agents is present in a sub-therapeutic dosage.

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16. A pharmaceutical composition according to Claim 15, wherein both the first and second therapeutic agents are present in sub-therapeutic dosages.

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17. A pharmaceutical composition according to Claim 14, further comprising:

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(d) a third therapeutic agent selected from other anti-coagulant or coagulation inhibitory agents, anti-platelet or platelet inhibitory agents, thrombin inhibitors, thrombolytic agents, fibrinolytic agents, anti-arrythmic agents, and cholesterol/lipid lowering agents.

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18. A pharmaceutical composition according to Claim 17, wherein the third therapeutic agent is selected from aspirin and pravastatin.

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19. A pharmaceutical composition according to Claim 18, wherein the third therapeutic agent is aspirin.

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20. A pharmaceutical composition according to Claim 17, wherein the third therapeutic agent is present in a sub-therapeutic dosage.

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